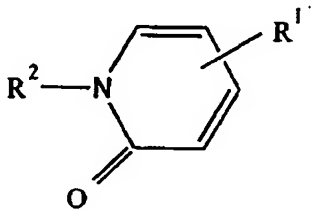
AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

LISTING OF CLAIMS:

1. (currently amended): A pharmaceutical liquid composition comprising ~~as an active ingredient~~ a pyridone derivative represented by the following formula (I):



wherein R^1 is an alkyl group optionally having a substituent selected from the group consisting of a C_{1-6} lower alkyl group optionally substituted at any of the 3-, 4- or 5-position with a halogen atom, a carboxyl group, an alkoxycarbonyl group, and an amino group and R^2 is a phenyl group optionally having a substituent selected from the group consisting of a C_{1-6} lower alkyl group, a halogen atom, a carboxyl group, an alkoxycarbonyl group or an amino group, or a pharmaceutically acceptable salt thereof, and a solvent capable of dissolving said pyridone derivative~~active ingredient~~ in a ~~high~~ concentration of about 10% to about 25% by weight.

2. (original): A pharmaceutical liquid composition according to Claim 1, ~~comprising as the active ingredient~~ wherein the pyridone derivative is a 5-methyl-1-phenyl-2-(1H)-pyridone (Pirfenidone) wherein R^1 is a methyl group at the 5-position and R^2 is a phenyl group in the formula (I) or a pharmaceutically acceptable salt thereof.

3. (currently amended): A pharmaceutical liquid composition according to Claim 1 ~~or 2~~, wherein the solvent is a diethylene glycol monoethyl ether.

4. (original): A pharmaceutical liquid composition according to Claim 3, wherein the diethylene glycol monoethyl ether has a purity of 99% or higher.

5. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to~~ 4, further comprising a concentrating agent.

6. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to~~ 5, further containing an antioxidant.

7. (original): A pharmaceutical liquid composition according to Claim 6, wherein the antioxidant is an α-tocopherol.

8. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to~~ 7, ~~which is suitable to be administered~~ in the form of an oral~~ly~~, percutaneous~~ly~~, nasal~~ly~~ or vaginal~~ly~~ preparation or ~~as~~ in the form of a spray, patch, inhalant, injection or intravenous drip.

9. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to~~ 8, having the following components:

Ingredients	% by weight
Pirfenidone	1-25

Diethylene glycol monoethyl ether	70-80
Ethanol (95%)	0-10
Polyvinyl pyrrolidone or hydroxypropyl cellulose	0-3
Sodium metabisulfite	0.02-2
Methyl or propyl paraben	0-0.5
Purified water	0-25 .

10. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to 8~~, having the following components:

Ingredients	% by weight
Pirfenidone	10-25
Diethylene glycol monoethyl ether	75-80
Purified water	0-10 .

11. (currently amended): A pharmaceutical liquid composition according to ~~any one of~~ Claims 1 ~~to~~ 8, having the following components:

Ingredients	% by weight
Pirfenidone	10-25

Diethylene glycol monoethyl ether	75-80
α-Tocopherol	0.1-0.5
Hydroxypropyl cellulose	0-3
Purified water	0-10 .